PSJ3 Exhibit 172

Tuesday, April 7, 2009

TOPIC	SUMMARY/DECISIONS	ACTIONS
Legal Statement	 Antitrust issue discussed by J. Dean, Covington & Burling There were no objections to proceeding with the meeting given this discussion 	• N/A
Welcome Introductions	Introductions of all participants on the telephone and those in the meeting room.	• N/A
Define operating principles of the Working Group Agree on key issues to be addressed/evaluated by the Working Group Determine next steps	There was agreement with the meeting goals No additional meeting goals were identified	• N/A
Operating Principles of Working Group Agree on primary member organizations/structure One collaborative Industry Working Group (IWG) or other options Key stakeholders Other Industry members not included in the FDA directive	The group agreed to proceed with one IWG to address the FDA directive The meeting of ANDA holders scheduled for April 20 th was discussed; it was agreed that this meeting would be held as an ANDA sub-team and the results reported back to the IWG It was agreed that the working group should only include the 26 companies identified by FDA as invitees to the March 3 rd meeting Additional input from other industry and non-industry (e.g., vendors) interested parties as well as stakeholders will be elicited through appropriate sub-teams It was agreed that a Charter would be drafted by a sub-team and would include the following working principles: Agree to abide by the antitrust guidelines Agree that all members will have an equal voice Agree to strive for consensus for all decisions Agree that membership will include the 26 member companies as identified by FDA	Sub-team to draft IWG charter and provide draft for discussion at next face-to-face meeting

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TOPIC	SUMMARY/DECISIONS	ACTIONS
Pain Care Forum		
Pain Care Forum REMS Task Force Class Opioids REMS Recommendations to FDA	 W. Rowe provided an overview of the PCF REMS Task Force and highlighted the current draft of their recommendations document which is 75-80% complete (slides to be provided) It was noted by Covidien that, although the document is called a "consensus" document, there were several key areas in which there was not "substantial agreement" by all PCF members It was suggested by Covidien that the core document include only those areas where there was substantial agreement and the other differing opinions be included as an appendix W. Rowe stated that the following meetings were scheduled at FDA and there would be representation by various members of the PCF: April 29 – AMA, medical specialties May 4 – prescribers May 4 – prescribers May 5 – pt. advocacy groups including PCF May 5 – dispensers (pharmacists) A 6th meeting will be requested by the PCF to discuss their recommendations which they plan to have complete by the third week in April It was agreed that the IWG should contact FDA about participating as an observer in these various meetings so that we can incorporate this stakeholder input into our REMS proposal 	Sub-team to contact FDA to discuss participation as an observer in various stakeholder meetings.

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Tuesday, April 7, 2009

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Assess the directive given to Industry by FDA What additional information do we need? What additional issues does this Working Group need to address? Which elements can this Working Group address? Which elements are outside of the scope of this Working Group? Engagement of key stakeholders	 The following issues were identified as needing additional information/ vetting: How do we measure change to benefit? Regarding the requirement of experience, training and/or certification as specified in FDAAA, "what" experience, "what" training and "what" certification will be required? What elements do we, as Industry, have the ability to influence or control? What are the Objectives of the proposed REMS that will allow us to meet the FDA stated Goals? 	Three sub-teams were formed to begin discussing potential Elements to Assure Safe Use (prescribers, patients, dispensers) – see Appendix 1 for subteam membership Engagement of key stakeholders will be determined by the sub-teams
■ Identification of sub-teams	Initially the following sub-teams were identified: Elements to Assure Safe Use Regulatory (FDA Communication) Legal Finance Logistics/Administrative Policy (to evaluate issues outside of the REMS directive such as inclusion of IR products) Evaluation/Metrics ANDA Holders Stakeholders PR – messaging After further discussion, it was agreed that that the following subteams would be formed at this time: Operations Communications Elements to Assure Safe Use Prescribers Patients Dispensers	Membership of subteams and point person outlined in Attachment 1 Provide TBD names to E. Smith by COB 4/8/2009 Sub-teams to convene by teleconference by April 10 to begin work and plan for agenda items for next face-to-face meeting Communication sub-team lead (E. Ernst) to contact FDA (B. Rappaport or S. Hertz) by phone to provide update on IWG An e-mail will also be drafted and circulated to the IWG for review prior to sending to FDA (E. Ernst)

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TOPIC	SUMMARY/DECISIONS	ACTIONS
Plan for the next meeting of the Working Group Establishment of a Charter	 Next IWG teleconference will be held the week of April 13 ANDA holder sub-team will meet on April 20 and Covidien will provide a report of this meeting to the IWG the week of April 27 Next face-to-face meeting will be held the week of May 11 so that an update from the FDA stakeholder meetings can be discussed Purdue agreed to host next face-to-face meeting in May Draft charter to be remit of Operations sub-team 	Sub-teams to convene TC by April 10 Operations sub-team to schedule IWG TC for week of April 13 th and the next face-to-face meeting for the week of May 11 th
Wrap-up • Summary/next actions		Minutes to be prepared and distributed by E. Smith
Adjourn		

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ATTACHMENT 1 - List of Sub-teams

Sub-team: Operations

Purpose: Coordinate logistics for meetings, draft IWG charter, investigate outside project

management support, discuss need for budget

Membership: Craig Landau – Purdue (point person)

Jill Buckley – King Kishore Gopu – Teva

Mylan (TBD) J&J (TBD)

Sub-team: Communications

Purpose: Communicate with FDA regarding status of IWG and request participation in

FDA-sponsored stakeholder meetings

Membership: Elizabeth Ernst – Roxane (point person)

Martin Lessem – Ranbaxy Marsha Stanton – King Michael Kaufman – J&J Burt Rosen – Purdue Tara Chapman – Endo Arthur Ilse – Xanodyne John Lay – Vista

Nick Tantillo – Teva/Barr Andrea Miller – Mylan Lorri Scheussler – Hisamitsu Beth Brannan – Watson Lesley Zhu – Sandoz

Sub-team: Elements to Assure Safe Use

Purpose: Gather information and evaluate potential Elements of Safe Use

Prescribers

Dave Haddox - Purdue (point person)

Eric Davis – Mylan Mike Nelson – Neuromed Sid Schnoll – Roxane

J&J - TBD

Shabana Modan - Ranbaxy

Eric Smith – King Frank Yuen – Endo

<u>Patients</u>

Sid Schnoll - Roxane (point person)

Shabana Modan - Ranbaxy

J&J - TBD

Lori Scheussler – Hisamitsu Deb Myers – Roxane

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ATTACHMENT 1 - List of Sub-teams (cont'd)

<u>Dispensers</u>
point person – TBD
Tara Chapman – Endo
Kimberly France – Covidien
Gary Kosloski – Watson
Purdue - TBD
J&J – TBD
Mylan – TBD
Sandoz – TBD

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